



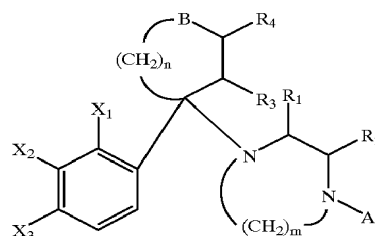
US005962455A

United States Patent [19]**Blum et al.**[11] **Patent Number:** **5,962,455**[45] **Date of Patent:** **Oct. 5, 1999**[54] **CERTAIN SUBSTITUTED BENZYLAMINE DERIVATIVES A NEW CLASS OF NEUROPEPTIDE Y1 SPECIFIC LIGANDS**[75] Inventors: **Charles A. Blum**, Guilford; **Alan Hutchison**, Madison; **John M. Peterson**, New Haven, all of Conn.[73] Assignee: **Neurogen Corporation**, Branford, Conn.[21] Appl. No.: **08/897,045**[22] Filed: **Jul. 18, 1997****Related U.S. Application Data**

[60] Provisional application No. 60/022,296, Jul. 23, 1996.

[51] **Int. Cl.⁶** **A61K 31/495**; C07D 295/135; C07D 401/12; C07D 403/10[52] **U.S. Cl.** **514/252**; 514/249; 514/253; 514/255; 540/470; 540/575; 544/356; 544/359; 544/295; 544/360; 544/362; 544/363; 544/370; 544/393[58] **Field of Search** 544/360, 363, 544/393, 295, 356, 359, 362, 370; 514/252, 253, 255, 249[56] **References Cited****FOREIGN PATENT DOCUMENTS**96143307 5/1996 WIPO .
9640660 12/1996 WIPO .**OTHER PUBLICATIONS**Gehlert et al., *Exp. Opin. Invest. Drugs*, 6 (12), pp. 1827-1838, 1997.Grundemar et al., *TIPS* 15, pp. 153-159, 1994.*Primary Examiner*—Emily Bernhardt*Attorney, Agent, or Firm*—Ladas & Parry[57] **ABSTRACT**

This invention encompasses compounds of the formula



and the pharmaceutically acceptable salts thereof

wherein X₁, X₂, X₃ represent organic or inorganic substituents, n is 1, 2, or 3, m is 2, 3, or 4, R₁–R₄ are hydrogen or organic substituents, and B is nitrogen, carbon, sulfur or oxygen, useful in the diagnosis and treatment of feeding disorders such as obesity and bulimia and cardiovascular diseases such as essential hypertension and congestive heart failure due to the binding of these compounds to mammalian Neuropeptide Y1 receptors.

11 Claims, No Drawings

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**CERTAIN SUBSTITUTED BENZYLAMINE
DERIVATIVES A NEW CLASS OF
NEUROPEPTIDE Y1 SPECIFIC LIGANDS**

This application claims priority from Provisional Patent Application Ser. No. 60/022,296 filed July 23, 1996. 5

BACKGROUND OF THE INVENTION

1. Field of the Invention

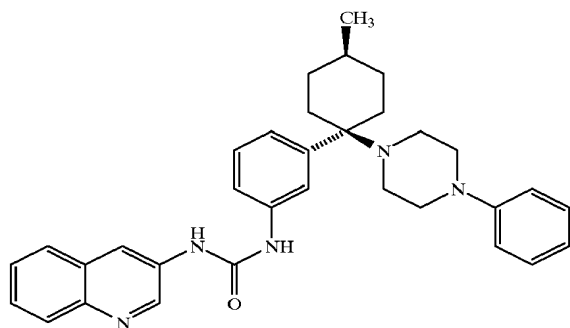
This invention relates to certain substituted benzylamine derivatives which selectively bind to mammalian Neuropeptide Y1 (NPY1) receptors. This invention also relates to pharmaceutical compositions comprising such compounds. It further relates to the use of such compounds and compositions in treating physiological disorders associated with an excess of Neuropeptide Y, especially feeding disorders and certain cardiovascular diseases.

2. Description of the Related Art

Neuropeptide Y, a peptide first isolated in 1982, is widely distributed in the central and peripheral neurons and is responsible for a multitude of biological effects in the brain and the periphery. Various animal studies have shown that activation of Neuropeptide Y1 receptors is related to vasoconstriction, Wahlestedt et al., Regul. Peptides, 13: 307-318 (1986), McCauley and Westfall, J. Pharmacol. Exp. Ther. 261: 863-868 (1992), and Grundemar et al., Br. J. Pharmacol. 105: 45-50 (1992); and to stimulation of consummatory behavior, Flood and Morley, Peptides, 10: 963-966 (1989), Leibowitz and Alexander, Peptides, 12: 1251-1260 (1991), and Stanley et al., Peptides, 13: 581-587 (1992).

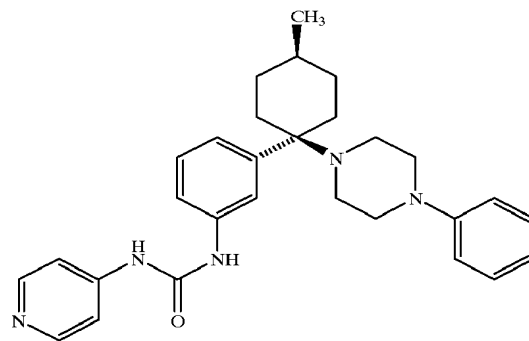
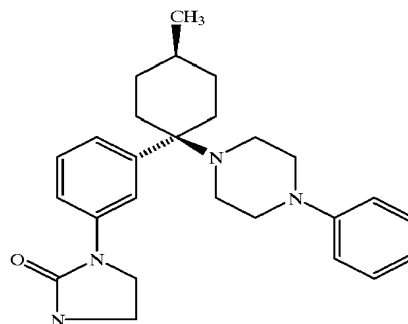
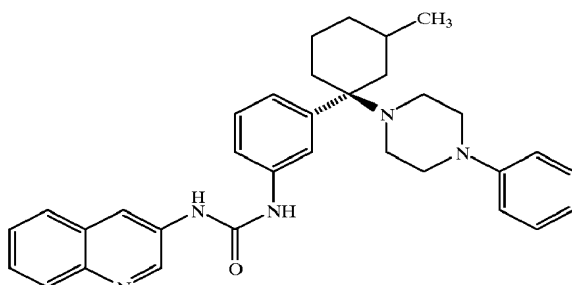
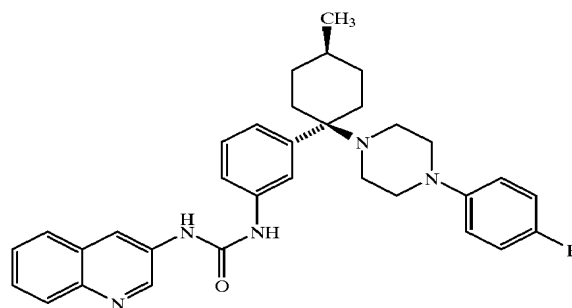
Grundemar and Hakanson, TIPS, May 1994 [Vol. 15], 153-159, state that, in animals, Neuropeptide Y is a powerful stimuli of food intake, and an inducer of vasoconstriction leading to hypertension. They further point out that low levels of Neuropeptide Y is associated with loss of appetite. These reports clearly indicate that compounds that inhibit the activity of this protein will reduce hypertension and appetite in animals.

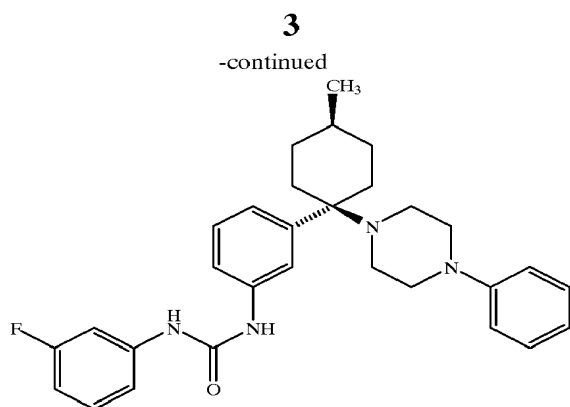
The following shows representative substituted benzylamines of the present invention.



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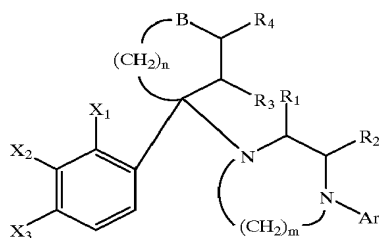




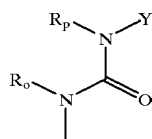
SUMMARY OF THE INVENTION

Compounds that interact with NPY1 receptors and inhibit the activity of Neuropeptide Y at those receptors are useful in treating physiological disorders associated with an excess of Neuropeptide Y such as eating disorders, for example, obesity and bulimia, and certain cardiovascular diseases, for example, hypertension.

The compounds encompassed by the instant invention are of the Formula I:



wherein one of X_1 , X_2 and X_3 is



and the remaining members of the group of X_1 , X_2 and X_3 are hydrogen; and where Y is an aryl group selected from the group consisting of phenyl, 2-, 3-, or 4-pyridyl, naphthyl, 2-, 3-, 4-, or 6-quinolyl, 3- or 4-isoquinolyl, 2- or 6-quinoxalyl, and 3-(1,8-naphthyridyl), each of which is optionally mono- or disubstituted with halogen, hydroxy, straight or branched chain C_1 - C_6 alkyl or C_1 - C_6 alkoxy;

R_0 and R_p are the same or different and represent hydrogen, straight or branched chain alkyl having 1-6 carbon atoms, aryl straight or branched chain lower alkyl having 1-6 carbon atoms or R_0 and R_p together may represent $-(CH_2)_n-$ where n is 1, 2 or 3; and

Ar is an aryl group preferably selected from the group consisting of phenyl, 2-, 3-, or 4-pyridyl, 2- or 3-thienyl, 2-, 4- or 5-pyrimidyl, each of which is optionally mono- or disubstituted with halogen, hydroxy, or straight or branched chain lower alkyl having 1-6 carbon atoms;

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B is sulfur, oxygen, $N(R_5)$ or $C(R_5)(R_6)$;

n is 1, 2, or 3;

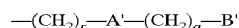
m is 2, 3, or 4;

R_1 and R_2 are the same or different and represent hydrogen, or straight or branched chain lower alkyl having 1-6 carbon atoms;

R_3 and R_4 are the same or different and represent hydrogen, straight or branched chain lower alkyl having 1-6 carbon atoms, or straight or branched chain lower alkoxy having 1-6 carbon atoms;

R_5 represents straight or branched chain lower-alkyl having 1-6 carbon atoms, phenyl, 2-, 3-, or 4-pyridyl, or phenyl, 2-, 3-, or 4-pyridyl straight or branched chain lower alkyl having 1-6 carbon atoms; and

R_6 represents hydrogen, hydroxyl, amino, straight or branched chain lower alkyl having 1-6 carbon atoms, straight or branched chain lower alkoxy having 1-6 carbon atoms, phenyl, 2-, 3-, or 4-pyridyl, phenoxy, 2-, 3-, or 4-pyridyloxy, or



where

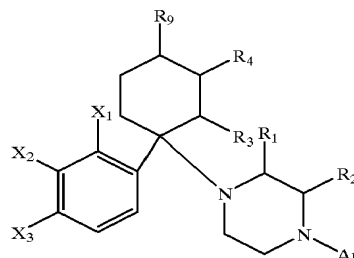
r is 0-5, q is 1-5, and A' is a direct bond, oxygen or sulfur, and

B' is hydrogen, straight or branched chain lower alkyl having 1-6 carbon atoms, straight or branched chain lower alkoxy having 1-6 carbon atoms, phenyl, 2-, 3-, or 4-pyridyl, phenoxy, 2-, 3-, or 4-pyridyloxy, carboxyl, carboalkoxy, carboxamido, mono or dialkylcarboxamido, amino, or mono or dialkylamino.

Preferred compounds according to Formula I are those where Ar is optionally substituted phenyl, pyrimidinyl or pyridyl, B is carbon optionally substituted with phenyl or alkyl, and R_1 - R_4 are hydrogen. Particularly, preferred compounds or Formula I are those where Ar is phenyl, pyrimidinyl or pyridyl, B is carbon optionally substituted with phenyl or alkyl, and R_1 - R_4 are hydrogen.

The invention also relates to compounds of formula IA:

IA

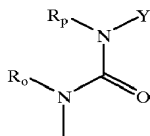


where

Ar is phenyl, 2-, 3-, or 4-pyridyl, 2- or 3-thienyl, 2-, 4- or 5-pyrimidyl, each of which is optionally mono- or disubstituted with halogen, hydroxy, straight or branched chain lower alkyl having 1-6 carbon atoms, or C_1 - C_6 alkoxy;

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wherein one of X_1 , X_2 or X_3 is



and the remaining members of the group of X_1 , X_2 and X_3 are hydrogen;

where Y is an aryl group preferably selected from the group consisting of phenyl, 2-, 3-, or 4-pyridyl, naphthyl, 2-, 3-, 4-, or 6-quinolyl, 3- or 4-isoquinolyl, 2- or 6-quinoxalyl, and 3-(1,8-naphthyridyl), each of which is optionally mono- or disubstituted with halogen, hydroxy, straight or branched chain lower alkyl having 1-6 carbon atoms, or C_1-C_6 alkoxy;

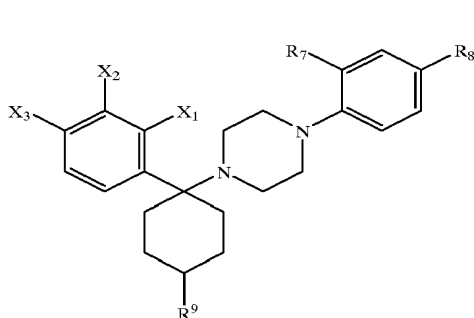
R_o and R_p are the same or different and represent hydrogen, straight or branched chain alkyl having 1-6 carbon atoms, aryl straight or branched chain lower alkyl having 1-6 carbon atoms or R_o and R_p together may represent $-(CH_2)_n$ where n is 1, 2 or 3; and

R_1 and R_2 are the same or different and represent hydrogen, or straight or branched chain lower alkyl having 1-6 carbon atoms;

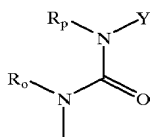
R_3 and R_4 are the same or different and represent hydrogen, straight or branched chain lower alkyl having 1-6 carbon atoms, or straight or branched chain lower alkoxy having 1-6 carbon atoms; and

R_9 represents hydrogen, straight or branched chain lower alkyl having 1-6 carbon atoms, phenyl.

The invention further encompasses compounds of Formula II:



where one of X_1 , X_2 and X_3 is



and the remaining members of the group of X_1 , X_2 and X_3 are hydrogen;

where Y is an aryl group preferably selected from the group consisting of phenyl, 2-, 3-, or 4-pyridyl, naphthyl, 2-, 3-, 4-, or 6-quinolyl, 3- or 4-isoquinolyl, 2- or 6-quinoxalyl, and 3-(1,8-naphthyridyl), each of which is optionally mono- or disubstituted with

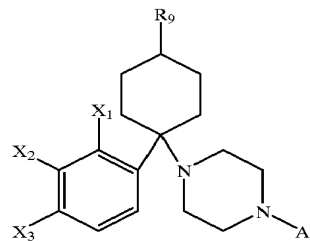
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halogen, hydroxy, straight or branched chain C_1-C_6 alkyl or C_1-C_6 alkoxy;

R_o and R_p are the same or different and represent hydrogen, straight or branched chain alkyl having 1-6 carbon atoms, aryl straight or branched chain lower alkyl having 1-6 carbon atoms or R_o and R_p together may represent $-(CH_2)_n$ where n is 1, 2 or 3; and where R_7 and R_8 are different and represent hydrogen or fluorine.

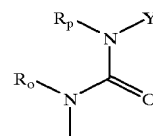
The invention also relates to compounds of Formula III:

III



where

Ar is phenyl, 2-, 3-, or 4-pyridyl, 2- or 3-thienyl 2-, 4- or 5-pyrimidyl, each of which is optionally mono- or disubstituted with halogen, hydroxy, or straight or branched chain lower alkyl having 1-6 carbon atoms; wherein one of X_1 , X_2 and X_3 is



and the remaining members of the group of X_1 , X_2 and X_3 are hydrogen;

where Y is an aryl group preferably selected from the group consisting of phenyl, 2-, 3-, or 4-pyridyl, naphthyl, 2-, 3-, 4-, or 6-quinolyl, 3- or 4-isoquinolyl, 2- or 6-quinoxalyl, and 3-(1,8-naphthyridyl), each of which is optionally mono- or distributed with halogen, hydroxy, straight or branched chain C_1-C_6 alkyl or C_1-C_6 alkoxy;

R_o and R_p are the same or different and represent hydrogen, straight or branched chain alkyl having 1-6 carbon atoms, aryl straight or branched chain lower alkyl having 1-6 carbon atoms or R_o and R_p together may represent $-(CH_2)_n$ where n is 1, 2 or 3; and

R_9 represents hydrogen, straight or branched chain lower alkyl having 1-6 carbon atoms, or phenyl.

The present invention includes compound of Formula I-III above and their pharmaceutically acceptable salts. Non-toxic pharmaceutically acceptable salts include salts of acids such as hydrochloric, phosphoric, hydrobromic, sulfuric, sulfonic, formic, toluene sulfonic, hydroiodic, acetic and the like. Those skilled in the art will recognize a wide variety of non-toxic pharmaceutically acceptable addition salts.

The present invention also encompasses the acylated prodrugs of the compounds of Formula I-III. Those skilled in the art will recognize various synthetic methodologies which may be employed to prepare non-toxic pharmaceuti-

cally acceptable addition salts and acylated prodrugs of the compounds encompassed by Formula I.

The invention encompasses both diastereomers of the compounds having 1,4-substitution on the cyclohexane ring, i.e. the invention encompasses both *cis*-, and *trans*-1,4-cyclohexanes. Preferred compounds of the invention having 1,4-substitution on the cyclohexane ring are those where the nitrogen atom forming the piperazine ring and the alkyl or phenyl group in the 4-position of the cyclohexane ring are “*cis*” with respect to each other. Thus, preferred compounds of the invention having such substitution are those that are *cis*-1-piperazinyl-4-alkyl or phenyl-cyclohexanes.

DETAILED DESCRIPTION OF THE INVENTION

By “aryl” and “Ar” is meant an aromatic carbocyclic group having a single ring (e.g., phenyl), multiple rings (e.g., biphenyl), or multiple condensed rings in which at least one is aromatic, (e.g., 1,2,3,4-tetrahydronaphthyl, naphthyl, anthryl, or phenanthryl), which can optionally be unsubstituted or substituted with e.g., halogen, lower alkyl, lower alkoxy, lower alkylthio, trifluoromethyl, lower acyloxy, aryl, heteroaryl, and hydroxy.

By “alkyl” and “lower alkyl” is meant straight and branched chain alkyl groups having from 1–6 carbon atoms.

By “lower alkoxy” and “alkoxy” is meant straight and branched chain alkoxy groups having from 1–6 carbon atoms.

By “halogen” is meant fluorine, chlorine, bromine and iodine.

As the compounds of Formula I are effective Neuropeptide Y1 receptor antagonists, these compounds are of value in the treatment of a wide variety of clinical conditions which are characterized by the presence of an excess of Neuropeptide Y. Thus, the invention provides methods for the treatment or prevention of a physiological disorder associated with an excess of Neuropeptide Y, which method comprises administering to a mammal in need of said treatment an effective amount of a compound of Formula I or a pharmaceutically acceptable salt, solvate or prodrug thereof. The term “physiological disorder associated with an excess of Neuropeptide Y” encompasses those disorders associated with an inappropriate stimulation of neuropeptide Y receptors, regardless of the actual amount of Neuropeptide Y present in the locale.

These physiological disorders may include:

- disorders or diseases pertaining to the heart, blood vessels or the renal system, such as vasospasm, heart failure, shock, cardiac hypertrophy increased blood pressure, angina, myocardial infarction, sudden cardiac death, arrhythmia, peripheral vascular disease, and abnormal renal conditions such as impaired flow of fluid, abnormal mass transport, or renal failure;

- conditions related to increased sympathetic nerve activity for example, during or after coronary artery surgery, and operations and surgery in the gastrointestinal tract;

- cerebral diseases and diseases related to the central nervous system, such as cerebral infarction, neurodegeneration, epilepsy, stroke, and conditions related to stroke, cerebral vasospasm and hemorrhage, depression, anxiety, schizophrenia, and dementia;

- conditions related to pain or nociception;

- diseases related to abnormal gastrointestinal motility and secretion, such as different forms of ileus, urinary incontinence, and Crohn’s disease;

- abnormal drink and food intake disorders, such as obesity, anorexia, bulimia, and metabolic disorders;
- diseases related to sexual dysfunction and reproductive disorders;

- conditions or disorders associated with inflammation; respiratory diseases, such as asthma and conditions related to asthma and bronchoconstriction; and diseases related to abnormal hormone release, such as leutinizing hormone, growth hormone, insulin, and prolactin. See U.S. Pat. No. 5,504,094.

The pharmaceutical utility of compounds of this invention is indicated by the following assay for human NPY1 receptor activity.

Assay for Human NPY1 Receptor Binding Activity

Membrane Preparation: Baculovirus-infected Sf9 cells expressing recombinant human NPY Y1 receptors were harvested at 42–48 hours at which time batches of 500 mL of cell suspension were pelleted by centrifugation. Each pellet was resuspended in 30 mL of lysis buffer (10 mM HEPES, 250 mM sucrose, 0.5 μ g/ml leupeptin, 2 μ g/ml Aprotinin, 200 μ M PMSF and 2.5 mM EDTA, pH 7.4) and gently homogenized by 50 strokes using a dounce homogenizer. The homogenate was centrifuged at 4° C. for 10 minutes at 536xg to pellet the nuclei. The supernatant was collected into a fresh tube and centrifuged twice in the same buffer at 48,000xg for 40 minutes. The final pellet was resuspended in 10 mL of PBS containing 5 mM EDTA by dounce homogenization and stored in aliquots at –80° C.

[¹²⁵I]PYY Binding Assay: Purified membranes were washed by PBS and resuspended by gentle pipetting in binding buffer [50 mM Tris(HCl), 5 mM KCl, 120 mM NaCl, 2 mM CaCl₂, 1 mM MgCl₂, 0.1% bovine serum albumin (BSA), pH 7.4]. Membranes (5 μ g) were added to siliconized (Sigma) polypropylene tubes in addition to 0.050 nM [¹²⁵I]PYY (porcine) for competition analysis or 0.010–0.500 nM [¹²⁵I]PYY (porcine) for saturation analysis. For evaluation of guanine nucleotide effects on receptor affinity, GTP was added at a final concentration of 100 μ M. Cold displacers were added at concentrations ranging from 10^{–12} M to 10^{–6} M to yield a final volume of 0.250 mL. Nonspecific binding was determined in the presence of 1 μ M NPY (human) and accounted for less than 10% of total binding. Following a 2 hour incubation at room temperature, the reaction was terminated by rapid vacuum filtration. Samples were filtered over presoaked GF/C Whatman filters (1.0% polyethylenimine for 2 hours) and rinsed 2 times with 5 mLs cold binding buffer lacking BSA. Remaining bound radioactivity was measured by gamma counting. To estimate the Bmax, Kd and Ki, the results of binding experiments were analyzed using SigmaPlot software (Jandel).

The compounds of general Formula I may be administered orally, topically, parenterally, by inhalation or spray or rectally in dosage unit formulations containing conventional non-toxic pharmaceutically acceptable carriers, adjuvants and vehicle. The term parenteral as used herein includes subcutaneous injections, intravenous, intramuscular, intrasternal injection or infusion techniques. In addition, there is provided a pharmaceutical formulation comprising a compound of general Formula I and a pharmaceutically acceptable carrier. One or more compounds of general Formula I may be present in association with one or more non-toxic pharmaceutically acceptable carriers and/or diluents and/or adjuvants and if desired other active ingredients. The pharmaceutical compositions containing compounds of general Formula I may be in a form suitable for oral use, for example, as tablets, troches, lozenges, aqueous or oily

suspensions, dispersible powders or granules, emulsion, hard or soft capsules, or syrups or elixirs.

Compositions intended for oral use may be prepared according to any method known to the art for the manufacture of pharmaceutical compositions and such compositions may contain one or more agents selected from the group consisting of sweetening agents, flavoring agents, coloring agents and preserving agents in order to provide pharmaceutically elegant and palatable preparations. Tablets contain the active ingredient in admixture with non-toxic pharmaceutically acceptable excipients which are suitable for the manufacture of tablets. These excipients may be for example, inert diluents, such as calcium carbonate, sodium carbonate, lactose, calcium phosphate or sodium phosphate; granulating and disintegrating agents, for example, corn starch, or alginic acid; binding agents, for example starch, gelatin or acacia, and lubricating agents, for example magnesium stearate, stearic acid or talc. The tablets may be uncoated or they may be coated by known techniques to delay disintegration and absorption in the gastrointestinal tract and thereby provide a sustained action over a longer period. For example, a time delay material such as glyceryl monostearate or glyceryl distearate may be employed.

Formulations for oral use may also be presented as hard gelatin capsules wherein the active ingredient is mixed with an inert solid diluent, for example, calcium carbonate, calcium phosphate or kaolin, or as soft gelatin capsules wherein the active ingredient is mixed with water or an oil medium, for example peanut oil, liquid paraffin or olive oil.

Aqueous suspensions contain the active materials in admixture with excipients suitable for the manufacture of aqueous suspensions. Such excipients are suspending agents, for example sodium carboxymethylcellulose, methylcellulose, hydropropylmethylcellulose, sodium alginate, polyvinylpyrrolidone, gum tragacanth and gum acacia; dispersing or wetting agents may be a naturally-occurring phosphatide, for example, lecithin, or condensation products of an alkylene oxide with fatty acids, for example polyoxyethylene stearate, or condensation products of ethylene oxide with long chain aliphatic alcohols, for example heptadecaethyleneoxycetanol, or condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with partial esters derived from fatty acids and hexitol anhydrides, for example polyethylene sorbitan monooleate. The aqueous suspensions may also contain one or more preservatives, for example ethyl, or n-propyl p-hydroxybenzoate, one or more coloring agents one or more flavoring agents, and one or more sweetening agents, such as sucrose or saccharin.

Oily suspensions may be formulated by suspending the active ingredients in a vegetable oil, for example arachis oil, olive oil, sesame oil or coconut oil, or in a mineral oil such as liquid paraffin. The oily suspensions may contain a thickening agent, for example beeswax, hard paraffin or cetyl alcohol. Sweetening agents such as those set forth above, and flavoring agents may be added to provide palatable oral preparations. These compositions may be preserved by the addition of an antioxidant such as ascorbic acid.

Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water provide the active ingredient in admixture with a dispersing or wetting agent, suspending agent and one or more preservatives. Suitable dispersing or wetting agents and suspending agents are exemplified by those already mentioned above.

Additional excipients, for example sweetening, flavoring and coloring agents, may also be present.

Pharmaceutical compositions of the invention may also be in the form of oil-in-water emulsions. The oily phase may be a vegetable oil, for example olive oil or arachis oil, or a mineral oil, for example liquid paraffin or mixtures of these. Suitable emulsifying agents may be naturally-occurring gums, for example gum acacia or gum tragacanth, naturally-occurring phosphatides, for example soy bean, lecithin, and esters or partial esters derived from fatty acids and hexitol anhydrides, for example sorbitan monooleate, and condensation products of the said partial esters with ethylene oxide, for example polyoxyethylene sorbitan monooleate. The emulsions may also contain sweetening and flavoring agents.

Syrups and elixirs may be formulated with sweetening agents, for example glycerol, propylene glycol, sorbitol or sucrose. Such formulations may also contain a demulcent, a preservative and flavoring and coloring agents. The pharmaceutical compositions may be in the form of a sterile injectable aqueous or oleaginous suspension. This suspension may be formulated according to the known art using those suitable dispersing or wetting agents and suspending agents which have been mentioned above. The sterile injectable preparation may also be sterile injectable solution or suspension in a non-toxic parentally acceptable diluent or solvent, for example as a solution in 1,3-butanediol. Among the acceptable vehicles and solvents that may be employed are water, Ringer's solution and isotonic sodium chloride solution. In addition, sterile, fixed oils are conventionally employed as a solvent or suspending medium. For this purpose any bland fixed oil may be employed including synthetic mono- or diglycerides. In addition, fatty acids such as oleic acid find use in the preparation of injectables.

The compounds of general Formula I may also be administered in the form of suppositories for rectal administration of the drug. These compositions can be prepared by mixing the drug with a suitable non-irritating excipient which is solid at ordinary temperatures but liquid at the rectal temperature and will therefore melt in the rectum to release the drug. Such materials are cocoa butter and polyethylene glycols.

Compounds of general Formula I may be administered parenterally in a sterile medium. The drug, depending on the vehicle and concentration used, can either be suspended or dissolved in the vehicle. Advantageously, adjuvants such as local anesthetics, preservatives and buffering agents can be dissolved in the vehicle.

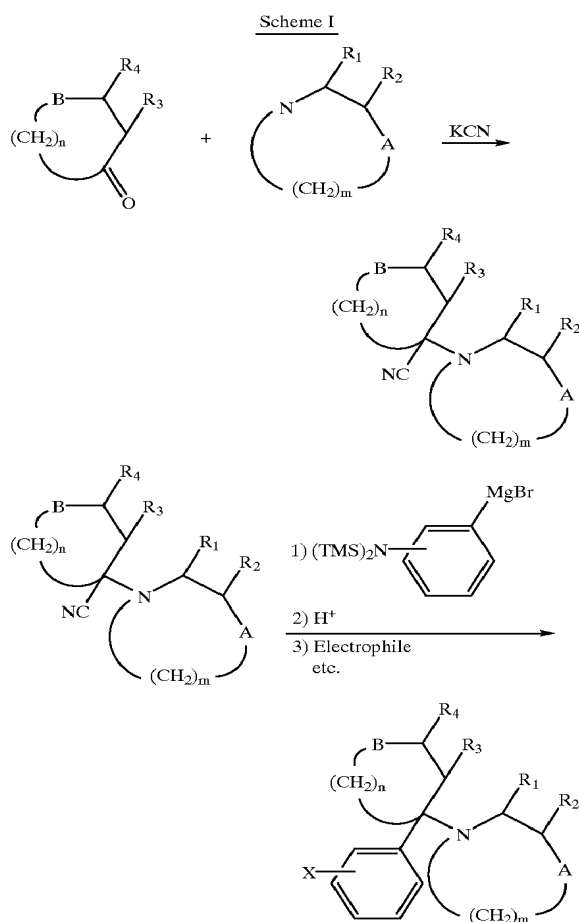
Dosage levels of the order of from about 0.1 mg to about 140 mg per kilogram of body weight per day are useful in the treatment of the above-indicated conditions (about 0.5 mg to about 7 g per patient per day). The amount of active ingredient that may be combined with the carrier materials to produce a single dosage form will vary depending upon the host treated and the particular mode of administration. Dosage unit forms will generally contain between from about 1 mg to about 500 mg of an active ingredient.

It will be understood, however, that the specific dose level for any particular patient will depend upon a variety of factors including the activity of the specific compound employed, the age, body weight, general health, sex, diet, time of administration, route of administration, and rate of excretion, drug combination and the severity of the particular disease undergoing therapy.

An illustration of the preparation of compounds of the present invention is given in Scheme I. Those having skill in the art will recognize that the starting materials may be varied and additional steps employed to produce compounds encompassed by the present invention.

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Starting materials are commercially available or may be prepared by procedures known to chemists of ordinary skill.



where

A is ArN or ArCH where Ar is phenyl, 2, 3, or 4 pyridyl, 2 or 3 thienyl, 2, 4 or 5 pyrimidyl either unsubstituted or mono or disubstituted with halogen, hydroxy, or straight or branched chain lower alkyl having 1-6 carbon atoms;

B is sulfur, oxygen NR₅ or CR₅R₆

n is 1, 2, or 3;

m is 2, 3, or 4;

R₁ and R₂ are the same or different and represent hydrogen, or straight or branched chain lower alkyl having 1-6 carbon atoms;

R₃ and R₄ are the same or different and represent hydrogen, straight or branched chain lower alkyl having 1-6 carbon atoms, or straight or branched chain lower alkoxy having 1-6 carbon atoms;

R₅ represents straight or branched chain lower alkyl having 1-6 carbon atoms, phenyl, 2, 3, or 4 pyridyl, or phenyl, 2, 3, or 4 pyridyl straight or branched chain lower alkyl having 1-6 carbon atoms;

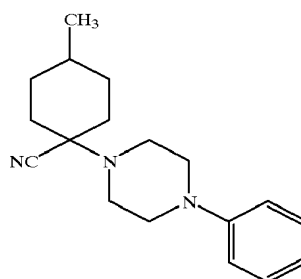
R₆ represents hydrogen, hydroxyl, amino, straight or branched chain lower alkyl having 1-6 carbon atoms, straight or branched chain lower alkoxy having 1-6 carbon atoms, phenyl, 2, 3, or 4 pyridyl, phenyloxy, 2, 3, or 4 pyridyloxy, or —(CH₂)_r—A'—(CH₂)_q—B'

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where r represents 0-5 and q represents 1-5 and A' is a direct bond, oxygen or sulfur and B' is hydrogen, straight or branched chain lower alkyl having 1-6 carbon atoms, straight or branched chain lower alkoxy having 1-6 carbon atoms, phenyl, 2, 3, or 4 pyridyl, phenyloxy, 2, 3, or 4 pyridyloxy, carboxyl, carboalkoxy, unsubstituted, mono or dialkylcarboxamido, amino, or mono or dialkylamino.

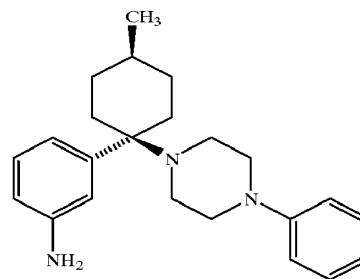
The invention is illustrated further by the following examples which are not to be construed as limiting the invention in scope or spirit to the specific procedures and compounds described in them.

EXAMPLE I



N-Phenylpiperazine (37 mL, 40 g, 245 mmol) was suspended in 300 mL water. The pH was adjusted to between 3 and 4 using 10% HCl. 4-Methyl cyclohexanone (30 mL, 27 g, 244 mmol) was added followed by KCN (16 g, 245 mmol). The mixture was stirred 15 hours at room temperature during which time the product solidified. The product was collected by filtration, washed with water, then dried in the vacuum oven overnight at 50° C. to give 58 g (84% yield) desired product as a roughly 2:1 mixture of diastereomers. Tic Rf=0.25 and 0.3 (9:1, Hexanes/Ethyl Acetate).

EXAMPLE II

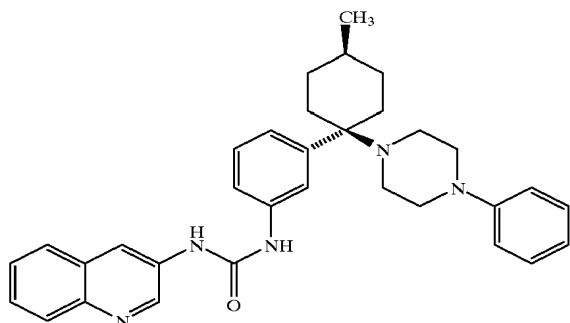


A 1 Molar THF solution of 3 [Bis(trimethylsilyl)amino] phenylmagnesium chloride (100 mL, 0.1 mol) was added to a solution of 1-cyano-1-(4-phenylpiperazine-1-yl)-4-methylcyclohexane (10 g, 0.035 mol) in dry THF (100 mL). The reaction mixture was heated to 65° C. for 2 h, cooled to room temperature and quenched by dropwise addition of saturated NH₄Cl solution. The magnesium salts were filtered, rinsed with THF and the filtrate was concentrated under reduced pressure. The residue was dissolved in EtOH (70 mL), 5% HCl solution (20 mL) was added, and the mixture stirred for 30 min. at room temperature. The mixture was filtered and the filtrate was concentrated under reduced pressure. The residue was suspended in H₂O, made basic

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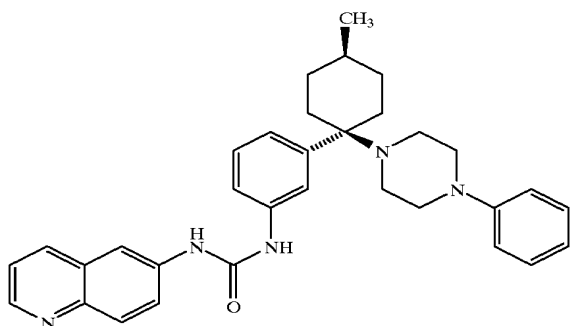
with 10 N NaOH and then extracted with EtOAc (3×). The combined extracts were washed with H₂O (1×) and brine (1×), dried (Na₂SO₄), filtered and concentrated under reduced pressure. The residue was filtered through silica gel (1:4/EtOAc:hexanes) and concentrated to give a pale yellow solid. Recrystallization from isopropyl alcohol yielded white needles of 1-(3-aminophenyl)-1-(4-phenylpiperazine-1-yl)-4-methyl-cyclohexane (cis isomer) in 38% yield. mp=142–144° C.

EXAMPLE III



A solution of 1-(3-aminophenyl)-1-(4-phenylpiperazin-1-yl)-4-methyl-cyclohexane (cis isomer, 90 mg, 0.258 mmol) and triethylamine (90 μ L, 0.65 mmol) in dry CH₂Cl₂ (7 mL) was brought to 0° C. using an ice bath. Phosgene (1.93 M in toluene, 160 μ L) was added dropwise and the resulting mixture was stirred at 0° C., under a dry N₂ atmosphere, for 30 minutes. 3-aminoquinoline (45 mg, 0.310 mmol) was added as a solid and the reaction was allowed to come to room temperature. Upon completion of the reaction (disappearance of 3-aminoquinoline) the mixture was diluted with an equal volume of CH₂Cl₂, washed with H₂O, dried (Na₂SO₄) and concentrated to give the crude urea. Silica gel chromatography yielded pure N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-3-quinoliny-urea (cis isomer) as an off-white solid. The HCl salt was prepared by adding excess saturated EtOAc/HCl solution to the free base in dry MeOH. Concentration of the homogeneous solution yielded a solid which was washed with EtOAc. mp=180–181° C.

EXAMPLE IV

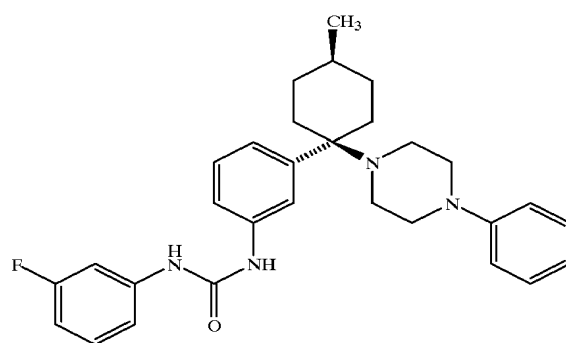


A solution of 1-(3-aminophenyl)-1-(4-phenylpiperazin-1-yl)-4-methyl-cyclohexane (cis isomer, 116 mg, 0.332 mmol) and triethylamine (116 μ L, 0.83 mmol) in dry CH₂Cl₂ (7

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mL) was brought to 0° C. using an ice bath. Phosgene (1.93 M in toluene, 181 μ L) was added dropwise and the resulting mixture was stirred at 0° C., under a dry N₂ atmosphere, for 30 minutes. 6-aminoquinoline (53 mg, 0.365 mmol) was added as a solid and the reaction was allowed to come to room temperature. The reaction was stirred overnight, diluted with an equal volume of CH₂Cl₂, washed with H₂O, dried (Na₂SO₄) and concentrated to give the crude urea. Silica gel chromatography (eluent: 5% CH₃OH/CH₂Cl₂) yielded pure N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-6-quinoliny-urea (cis isomer) as an off-white solid. The HCl salt was prepared by adding excess saturated EtOAc/HCl solution to the free base in dry MeOH. Concentration of the homogeneous solution yielded a white solid which was washed with EtOAc. m=185–187° C.

EXAMPLE V



A solution of 1-(3-aminophenyl)-1-(4-phenylpiperazin-1-yl)-4-methyl-cyclohexane (cis isomer, 205 mg, 0.587 mmol) and triethylamine (205 μ L, 1.47 mmol) in dry CH₂Cl₂ (15 mL) was brought to 0° C. using an ice bath. Phosgene (1.93 M in toluene, 365 μ L) was added dropwise and the resulting mixture was stirred at 0° C., under a dry N₂ atmosphere, for 30 minutes. 3-fluoroaniline (68 μ L, 0.705 mmol) was added via syringe and the reaction was allowed to come to room temperature. After 3 h the mixture was diluted with an equal volume of CH₂Cl₂, washed with H₂O, dried (Na₂SO₄) and concentrated to give the crude urea. The residue was triturated with ether and the resulting solid collected on a sintered glass funnel and washed with ether. Preparative plate chromatography (50% EtOAc/hexanes) afforded pure N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-3-fluorophenyl-urea (cis isomer) as an off-white solid. The HCl salt was prepared by adding excess saturated EtOAc/HCl solution to the free base in dry MeOH. Concentration of the homogeneous solution yielded a solid which was washed with EtOAc and dried under vacuum. mp=149–151° C.

EXAMPLE VI

The following compounds were prepared essentially according to the procedure described in Examples I–V:

- N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-2-pyridyl-urea trihydrochloride (cis isomer: Compound 4). mp=169–171° C.
- N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-3-pyridyl-urea trihydrochloride (cis isomer: Compound 5). mp=186–188° C.
- N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-2-naphthyl-urea dihydrochloride (cis isomer: Compound 6). mp=165–167° C.

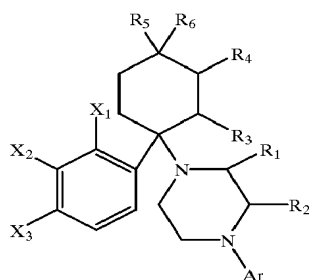
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- d) N-[3-[3-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-3-quinolinyl-urea trihydrochloride (trans isomer: Compound 7). mp=168–170° C.
- e) N-[3-[4-methyl-1-(4-(4-fluoro)phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-3-quinolinyl-urea trihydrochloride (cis isomer: Compound 8). mp=170–172° C.
- f) N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-4-fluorophenyl-urea dihydrochloride (cis isomer: Compound 9). mp=150–152° C.

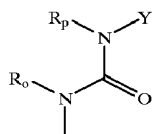
The invention and the manner and process of making and using it, are now described in such full, clear, concise and exact terms as to enable any person skilled in the art to which it pertains, to make and use the same. It is to be understood that the foregoing describes preferred embodiments of the present invention and that modifications may be made therein without departing from the spirit or scope of the present invention as set forth in the claims. To particularly point out and distinctly claim the subject matter regarded as invention, the following claims conclude this specification.

We claim:

1. A compound of the formula:



wherein one of X₁, X₂ and X₃ is



and the remaining members of the group of X₁, X₂ and X₃ are hydrogen;

where Y is an aryl group selected from the group consisting of phenyl, 2-, 3-, or 4-pyridyl, naphthyl, 2-, 3-, 4-, or 6-quinolyl, 3- or 4-isoquinolyl, 2- or 6-quinoxalyl, and 3-(1,8-naphthyridyl), each of which is optionally mono- or disubstituted with halogen, hydroxy, straight or branched chain C₁–C₆ alkyl or C₁–C₆ alkoxy;

R_o and R_p are the same or different and represent hydrogen-, straight or branched chain alkyl having 1–6 carbon atoms, arylalkyl wherein the alkyl portion is straight or branched chain lower alkyl having 1–6 carbon atoms or R_o and R_p together may represent —(CH₂)_n— where n is 1, 2 or 3; and

Ar is phenyl which is optionally mono- or disubstituted with halogen, hydroxy, or straight or branched chain lower alkyl having 1–6 carbon atoms; m is 2,

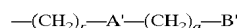
R₁ and R₂ are the same or different and represent hydrogen-, straight or branched chain lower alkyl having 1–6 carbon atoms;

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R₃ and R₄ are the same or different and represent hydrogen, straight or branched chain lower alkyl having 1–6 carbon atoms, or straight or branched chain lower alkoxy having 1–6 carbon atoms;

R₅ represents straight or branched chain lower alkyl having 1–6 carbon atoms, phenyl, 2-, 3-, or 4-pyridyl, or phenyl, 2-, 3-, or 4-pyridyl straight or branched chain lower alkyl having 1–6 carbon atoms; and

R₆ represents hydrogen, hydroxyl, amino, straight or branched chain lower alkyl having 1–6 carbon atoms, straight or branched chain lower alkoxy having 1–6 carbon atoms, phenyl, 2-, 3-, or 4-pyridyl, phenoxy, 2-, 3-, or 4-pyridyloxy, or

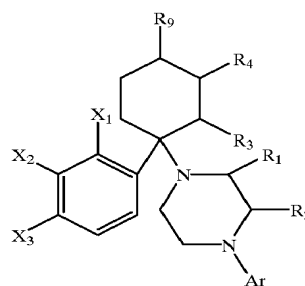


where

r is 0–5, q is 1–5, and A' is a direct bond, oxygen or sulfur, and

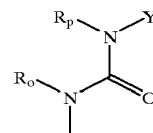
B' is hydrogen, straight or branched chain lower alkyl having 1–6 carbon atoms, straight or branched chain lower alkoxy having 1–6 carbon atoms, phenyl, 2-, 3-, or 4-pyridyl, phenoxy, 2-, 3-, or 4-pyridyloxy, carboxyl, carboalkoxy, carboxamido, mono or dialkylcarboxamido, amino, or mono or dialkylamino; and pharmaceutically acceptable salts thereof.

2. A compound of the formula:



where

Ar is phenyl which is optionally mono or disubstituted with halogen, hydroxy, straight or branched chain lower alkyl having 1–6 carbon atoms or C₁–C₆ alkoxy; one of X₁, X₂, or X₃ is



and the remaining members of the group X₁, X₂ and X₃ are hydrogen;

Y is an aryl group selected from the group consisting of phenyl, 2-, 3-, or 4-pyridyl, naphthyl, 2-, 3-, 4-, or 6-quinolyl, 3- or 4-isoquinolyl, 2- or 6-quinoxalyl, and 3-(1,8-naphthyridyl), each of which is optionally mono- or disubstituted with halogen, hydroxy, straight or branched chain C₁–C₆ alkyl or C₁–C₆ alkoxy;

R_o and R_p are the same or different and represent hydrogen-, straight or branched chain alkyl having 1–6 carbon atoms, arylalkyl wherein the alkyl portion is straight or branched chain lower alkyl having 1–6

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carbon atoms or R_o and R_p together may represent $-(CH_2)_n-$, where n is 1, 2 or 3; R_1 and R_2 are the same or different and represent hydrogen, or straight or branched chain lower alkyl having 1–6 carbon atoms;

R_3 and R_4 are the same or different and represent hydrogen, straight or branched chain lower alkyl having 1–6 carbon atoms, or straight or branched chain lower alkoxy having 1–6 carbon atoms; and

R_5 represents hydrogen, straight or branched chain lower alkyl having 1–6 carbon atoms or phenyl.

3. A compound as claimed in claim 1 wherein Ar is phenyl optionally mono- or disubstituted with halogen, hydroxy or straight or branched chain lower alkyl having 1–6 carbon atoms.

4. A compound as claimed in claim 1 wherein Ar is phenyl.

5. A compound according to claim 1, which is N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-2-pyridyl-urea trihydrochloride (cis isomer), N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-3-pyridyl-urea trihydrochloride (cis isomer), N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-2-naphthyl-urea dihydrochloride (cis isomer), N-[3-[3-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-3-quinoliny-urea trihydrochloride (trans isomer), N-[3-[4-methyl-1-(4-(4-fluoro)phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-3-quinoliny-urea trihydrochloride (cis isomer), N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl) cyclohexyl]

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phenyl]-N'-4-fluorophenyl-urea dihydrochloride (cis isomer), N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-3-quinoliny-urea trihydrochloride (cis isomer), N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-6-quinoliny-urea trihydrochloride (cis isomer) or N-[3-[4-methyl-1-(4-phenyl-1-piperazinyl)cyclohexyl]phenyl]-N'-3-fluorophenyl-urea dihydrochloride (cis isomer).

6. A method of treating an eating disorder in a mammal which comprises administering to a mammal in need of such treatment an effective amount of a compound as claimed in claim 1.

7. A pharmaceutical composition comprising a compound of claim 1, and a pharmaceutically acceptable carrier.

8. A method of treating an eating disorder in a mammal which comprises administering to a mammal in need of such treatment an effective amount of a compound as claimed in claim 2.

9. A method of treating an eating disorder in a mammal which comprises administering to a mammal in need of such treatment an effective amount of a compound as claimed in claim 5.

10. A pharmaceutical composition comprising a compound of claim 2 and a pharmaceutically acceptable carrier.

11. A pharmaceutical composition comprising a compound of claim 5 and a pharmaceutically acceptable carrier.

* * * * *